

L 51980-65

ACCESSION NR: AT5012684

tungsten from solution in a single operation under optimum conditions. It was postulated that tungsten penetrates the anions $[PMo_{12}O_{40}]^{3-}$ to form the poly-anions $[PW_nMo_{12-n}O_{40}]^{3-}$. A technique was worked out for determining 1×10^{-4} to $2 \times 10^{-2}\%$ tungsten in molybdenum. The coprecipitation of 0.01-1.0 mg of Nb and Ta with ammonium phosphomolybdate was also studied. A spectrochemical method was used to follow the coprecipitation of Nb and Ta with other ions such as aluminum, iron, and titanium, whose total oxides are present in amounts 10^4 - 10^5 times greater than Ta and Nb. The study led to the development of a technique for determining these two elements in rocks with large amounts of sesquioxides. A method was also proposed for a secondary concentration of tungsten, niobium, and tantalum by the vacuum distillation of $P_2O_5 \cdot xMoO_3$. Orig. art. has: 6 figures and 2 tables.

ASSOCIATION: Komissiya po analiticheskoy khimii, AN SSSR (Commission for Analytical Chemistry, AN SSSR)

SUBMITTED: 00

ENCL: 00

SUB CODE: IC, MM

NO REF SOV: 006

OTHER: 000

ml
Card 2/2

SEMENENKO, K. N.

Basic beryllium acetate. I. Monoclinic high temperature modification of basic beryllium acetate. K. N. Semenenko, Yu. P. Simanov, and A. V. Novoselova. *Vestnik Dneprov. Univ.*, No. 2; Ser. Fiz.-Mat. i Estestv. Nauk No. 1, 61-2 (1954). Basic Be acetate prepd. either by sublimation, crystals of the melt; or crystals from BuOH; forms monoclinic crystals with cell dimensions: a 13.6; b 9.24; c 16.20 Å, β 80°. Pycnometric d. 1.340. A powder and heated (180-6°) specimen shows d. 1.335 which gives λ for a monoclinic lattice of 4.03, or λ for a rhombohedral lattice of 1.39. Thus the previously reported transition of the cubic to the rhombohedral form at 148° is in error (S. Seki, *et al.*, *Nature*, 163, 225 (1949)). G. Kosolapoff.

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Chin. Inorg. Chem.

SEMENENKO, K. N.

USSR/Chemistry

FD-775

Card 1/1 : Pub 129 12/24

Author : Lapitskiy, A. V.; Simanov, Yu. P.; Semenenko, K. N.; Yarembash, Ye. I.

Title : Some properties of tantalum pentoxide

Periodical : Vest. Mosk. un., Ser. fizikomat. i yest. nauk, Vol 9, No 2, 85-89,
Mar 1954

Abstract : Studied the dehydration process of tantalum pentoxide hydrate in the temperature range of 25-450 degrees. Established the possibility of the existence of a tantalic acid with the composition $H_7 [Ta(TaO_4)_4]$. Also studied the possible polymorphic conversions of tantalum pentoxide using X-ray and thermographic techniques. Determined the parameters of two modifications of tantalum pentoxide indicated in the rhombic lattice. Expressed an assumption regarding the possibility of the existence of a tantalic acid having the composition $H_{13} [Ta (TaO_4)_6]$. One table. Eight references (three foreign).

Institution : Chair of Inorganic Chemistry

Submitted : July 11, 1953

FD-2167

USSR/Chemistry - Inorganic

Card 1/1 Pub 129-7/20

Author : Novoselova, A. V.; Pashinkin, A. S.; Semenenko, K. N.

Title : Investigating the system NH_4Cl - BeCl_2 by thermometric titration and solubility determination

Periodical : Vest. Mos. un., Ser. fizikomat. i yest. nauk, 10, No 2, 49-56, Mar 1955

Abstract : Confirmed the formation of complexes between beryllium and ammonium chlorides by thermometric titration (plotting temperature vs composition and using maxima as indications of complex formation.). Also obtained solubility data on the system NH_4Cl - BeCl_2 - H_2O . Tables, diagrams, Fourteen references (six USSR; eight since 1940).

Institution : Chair of Inorganic Chemistry

Submitted : September 2, 1954

NOVOSELOVA, A.V.; PASHINKIN, A.S.; SEMENENKO, K.N.; YAREMBASH, Ye.I.

Instrument designed for laboratory work with hygroscopic and
hydrolyzing substances. Zav.lab.21 no.7:857-858 '55.

(MLRA 8:10)

1. Moskovskiy gosudarstvennyy universitet
(Chemical apparatus)

SEMENENKO, K.N.

Compounds of beryllium oxycarboxylates with pyridine and dioxane. A.V. Novoselov, Yu. P. Smirnov, K. N. Semenenko, and N. N. Krasovskii. Zhur. Neorgan. Khim. 1: 141-146 (1956). A detailed investigation was made of the compounds of beryllium oxycarboxylates with pyridine, $\text{Be}_2\text{O}(\text{CH}_3\text{COO})_4 \cdot 3\text{C}_5\text{H}_5\text{N}$, and with dioxane, $\text{Be}_2\text{O}(\text{CH}_3\text{COO})_4 \cdot \text{C}_4\text{H}_8\text{O}_2$. These materials are insoluble in organic solvents but are readily soluble in water and water solution of both acids and bases. The process of thermal decomposition of the tripyridinate of beryllium oxycarboxylate was investigated. The data seem to show that the compound $\text{Be}_2\text{O}(\text{CH}_3\text{COO})_4 \cdot 3\text{C}_5\text{H}_5\text{N}$ is a solid solution of pyridine in the crystal lattice of an unstable molecular compound of composition $\text{Be}_2\text{O}(\text{CH}_3\text{COO})_4 \cdot \text{C}_5\text{H}_5\text{N}$. 7 figures. 1 reference.

U.S.A.

Semenko, R. N.

1
Beryllium oxymonochloroacetate (A. V. Novoselova and R. N. Semenko, *Zh. Neorg. Khim.* 1987, 14 (1964)).
The lattice const. of II were determined by x-ray diagrams and compared with those for I. The only significant difference is in the values of c ($c_1 = 0.15$ kX; $c_2 = 7.3$ kX). The difference in the mol. structure in the case of II is attributed to the strength of the acid ClCH_2COOH as compared to acids of the type RCOOH , where R is a hydrocarbon radical.
Rovnar, Leach

2

PM

SEMENENKO, K. N.

C.

USSR/Inorganic Chemistry - Complex Compounds.

Abs Jour : Ref Zhur - Khimiya, No 9, 1957, 30322

Author : Novoselova, A.V., Semenenko, K.N.

Inst : Interaction of Beryllium Oxyacetate with Beryllium

Title : Oxymonochloracetate.

Orig Pub : Zh. neorgan. khimii, 1956, 1, No 10, 2344-2348

Abst : By methods of thermal and x-ray phase analyses a study was made of the system $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_6$ (I) - $\text{Be}_4\text{O}(\text{CH}_2\text{ClCOO})_6$ (II). A chemical interaction takes place in the system between I and II, which results in the formation of four phases of variable composition which crystallize on the basis of I, II, $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_4(\text{CH}_2\text{ClCOO})_2$ (III) and $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_2(\text{CH}_2\text{ClCOO})_4$. The system is characterized by two eutectics: 72% I at 210° and 23% I at 190° , peritectic

Card 1/2

СЕМЕНЕНКО, К.Н.

TUROVA, N.Ya.; NOVOSELOVA, A.V.; SEMENENKO, K.N.

Reaction of beryllium hydroxyacetate with ammonium difluoride.

Zhur. neorg. khim. 1 no.11:2567-2569 N '56.

(MLRA 10:5)

1. Moskovskiy gosudarstvennyy universitet im. M.V. Lomonosova.
(Beryllium acetates) (Ammonium fluorides)

SEMENENKO, A. N.

C

USSR / Inorganic Chemistry. . Complex Compounds

Abs Jour : Ref Zhur - Khimiya, No 3, 1957, No 7797

Author : Novoselova, A.V., Semenenko, K.N., Krasovskaya, N.N. and Simanov, Yu.P.

Inst : Moscow University

Title : Beryllium Oxyacetate. Communication 2. Concerning Some Properties of Beryllium Oxyacetate-Pyridine Compounds

Orig Pub : Vestn. Mosk. Un-ta, 1956, No 3, 87-93

Abstract : Barium oxyformate, $\text{Be}_4(\text{HCOO})_6$ (I), has been synthesized and investigated and the formation and the properties of compounds of I, beryllium oxyacetate ($\text{Be}_4\text{O}(\text{CH}_3\text{COO})_6$ (II), and beryllium oxypropionate ($\text{Be}_4\text{O}(\text{CH}_2\text{CHCOO})_6$ (III) with pyridine (IV) and dioxene (V) have been studied. I was prepared by treating Be hydroxide or bicarbonate with formic acid, followed by the decomposition of the normal Be formate which is obtained in vacuo at $250 - 260^\circ$. At 250° , the yield of pure I

Card : 1/4

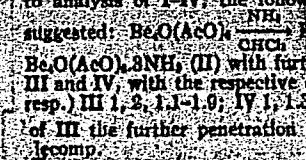
SEMENENKO, K. N.

Beryllium oxycacetate. III.

Novoselova, and K. N. Semenenko, *Vysok. Khim.*

Unio. Ser. Mat. Khim. Anal. 11, No. 2, 201-4 (1956); cf. C.A. 50, 163 (1956).

Be₂O(AcO)₄, dissolved in CHCl₃, produces a ppt. (I), quickly dissolving in excess of NH₃, and forming a product (II), but giving way to a heavy insol. final ppt. (III). The soln. contains a substance (IV), different from III. According to analysis of I-IV, the following scheme of reaction is suggested:

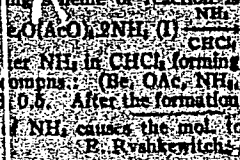


Interaction of beryllium oxycacetate with ammonia?

N. Maksimov, A. V. Semenenko, *Vysok. Khim.*

Unio. Ser. Mat. Khim. Anal. 11, No. 2, 201-4 (1956); cf. C.A. 50, 163 (1956).

Be₂O(AcO)₄, dissolved in CHCl₃, produces a ppt. (I), quickly dissolving in excess of NH₃, and forming a product (II), but giving way to a heavy insol. final ppt. (III). The soln. contains a substance (IV), different from III. According to analysis of I-IV, the following scheme of reaction is suggested:



SEMEENKO, K. N

USSR/Inorganic Chemistry. Complex Compounds.

C

Abs Jour: Ref. Zhur-Khimiya, No 1, 1958, 663.

Author : Maksimov, V.N., Novoselova, A.V., Semenenko, K.N.

Inst : -

Title : Lanthanum Acetate.

Orig Pub: Zh. Neorgan. Khimii, 1957, 2, No 5, 997-1000

Abstract: When $\text{La}(\text{CH}_3\text{COO})_3 \cdot 1.5 \text{H}_2\text{O}$ is heated at $50-110^\circ$, 0.5 molecule of water is lost and the monohydrate is formed which subsequently is completely dehydrated during heating at $110-150^\circ$. In the course of this treatment the dehydrated $\text{La}(\text{CH}_3\text{COO})_3$ is partially decomposed and at a temperature of $\sim 300^\circ$ it is converted into "metaacetate" $\text{LaO}(\text{CH}_3\text{COO})$. During boiling of the $\text{La}(\text{CH}_3\text{COO})_3 \cdot 1.5 \text{H}_2\text{O}$ in acetic anhydride a water-free lanthanum acetate is obtained that is stable in the air at room temperature. During heating to 300° it also is converted to $\text{LaO}(\text{CH}_3\text{COO})$.

Card : 1/1

-9-

SEMENENKO, K.N.

SEMENENKO, K.N.

Salts of zinc and cadmium with the simplest organic acids of the
type RCOOH. Zhur.neorg.khim. 2 no.9:2115-2121 S '57. (MIRA 10:12)
(Zinc salts) (Cadmium salts) (Acids, Organic)

SEMSNERO, K. N.

11str: 4E3j/4E3d/4E2c(1)

Some properties of barium perchlorate, $\text{Ba}(\text{ClO}_4)_2$, prepared by Kondyminov and K. N. Semner (State Univ. Moscow) (Vestnik Akad. Nauk, 12, Ser. Khim. No. 3, 205-7 (1957)).
 by adding $\text{Ba}(\text{OH})_2$ to a BaCO_3 - H_2O slurry, and extr. with CHCl_3 (40-50% yield). It was also prep. by adding II to melted $\text{Ba}(\text{OH})_2$, washing, and extr. (80% yield). Recryst. I. m. 115-116° shows endothermic effects at 200° (irreversible) and starts to sublime at 305°. Sublimation of I is stable above 200°. The low-temp. I is monoclinic with cell parameters of $a = 21.02$, $b = 11.1$, $c = 12.3$ Å. By absorbing 2-3 mole. of C_2H_4 the latter forms an unstable compound, which was prep. with I and other org. solvents.

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5(3)
AUTHORS: Semenenko, K.N. and Kurdyumov, G.M. SOV/55-58-2-28/35
TITLE: On Complex Compounds of Zinc Acetate With Nitrogenous Organic Substances (O kompleksnykh soyedineniyakh uksusnokislogo tsinka s azotsoderzhashchimi organicheskimi veshchestvami)
PERIODICAL: Vestnik Moskovskogo Universiteta. Seriya matematiki, mekhaniki, astronomii, fiziki, khimii, 1958, Nr 2, pp 207-210 (USSR)
ABSTRACT: The compounds $\text{Zn}(\text{CH}_3\text{COO})_2 \cdot 2\text{C}_5\text{H}_5\text{N}$ and $\text{Zn}(\text{CH}_3\text{COO})_2 \cdot 2\text{C}_4\text{H}_9\text{NH}_2$ were produced and investigated. It was shown that zinc acetate solutions in water and several organic solvents possess an abnormally high viscosity. The monocrystals of the produced combinations mentioned above were radiographically investigated. There are 3 tables, and 8 references, 1 of which is Soviet, 3 are German, 2 French, and 2 American.
SUBMITTED: June 7, 1957

Card 1/1

AUTHOR: Serenenko, K. N. SCV/156 58-2-19/48

TITLE: Concerning the Zinc Salts of Propionic and Butyric Acids
(O sol'yakh tsinka s propionovoy i maslyanoy kislota)

PERIODICAL: Nauchnyye doklady vysshey shkoly. Khimiya i khimicheskaya
tekhnologiya, 1958, Nr 2, pp. 283 - 284 (USSR)

ABSTRACT: Zinc propionate and zinc butyrate are only slightly soluble in
water or in aqueous solutions of propionic or butyric acid,
and they form no hydrated crystals. They can be converted to
zinc oxide by heating in hot concentrated acid. These salts
crystallize as small, very soft, thin transparent platelets.
These crystals were washed with absolute ether and analysed.
Zinc propionate and zinc butyrate are practically insoluble in
such solvents as benzene, ether, and chloroform. At 150-200°
and atmospheric pressure they sublime without decomposition.
Figure 1 shows the "thermogram" for heating each salt. The
endothermic effects at 205° and 160° (Fig 2, curve "4")
correspond to the melting points of these salts. In figure 2
these endothermic effects at the melting point are compared
to that for zinc acetate (non-aqueous). The linear relationship

Card 1/2

Concerning the Zinc Salts of Propionic and Butyric
Acids

SOV/236-58-2-19/48

between the melting temperatures of the salts of type $Zn(RCOO)_2$ and the number of carbon atoms in each salt is striking. The results of x-ray investigations on both salts are given. Because of their low melting points these salts probably do not exist in a crystalline lattice. They are remarkable for their relatively weak intermolecular bonds.

ASSOCIATION: Kafedra neorganicheskoy khimii Moskovskogo gosudarstvennogo universiteta im. M. V. Lomonosova (Chair of Inorganic Chemistry of the Moscow State University imeni M. V. Lomonosov)

SUBMITTED: November 5 1957

Card 2/2

5(2)
 AUTHORS: Semenenko, K.N. and Kurdyumov, G.M. SOV/55-58-3-22/30
 TITLE: On the Combinations of Beryllium Hydroxybenzoate With Toluene and Halogen-Substituted Benzene (O soyedineniyakh oksibenzoata berilliya s toluolom i galogenzameshchennymi benzolami)
 PERIODICAL: Vestnik Moskovskogo universiteta, Seriya matematiki, mekhanika, astronomii, fiziki, khimiya, 1958, Nr 3, pp 187-190 (USSR)
 ABSTRACT: By vaporization of a Be-hydroxybenzoate solution in a corresponding solvent there were obtained in crystalline form less stable combinations of Be-hydroxybenzoate with some organic molecules. A radiography was carried out and it showed that the organic molecules are enclosed in the interspaces of the crystal lattice of Be-hydroxybenzoate, whereby the original lattice is somewhat deformed. There are 2 tables, and 2 Soviet references.
 ASSOCIATION: Kafedra neorganicheskoy khimii (Chair of Inorganic Chemistry)
 SUBMITTED: June 7, 1957

Card 1/1

78-3-6-28/30

Author: Gerasimov, K. N.

Topic: On the Acetates of Zinc and Beryllium (Ob acetatakh tsinka i berilliya)

Periodical: Zhurnal Neorganicheskoy Khimii, 1958, Vol. 3, No. 6, pp. 1467-1468 (USSR)

Abstract: The syntheses of oxy-acetates and acetates of zinc and beryllium are described. The synthesis of oxy-acetate of zinc was obtained by the thermal decomposition of $\text{Zn}(\text{CH}_3\text{COO})_3 \cdot 2 \text{H}_2\text{O}$ in vacuum at 280°C . Zinc oxy-acetate is isomorphous with beryllium oxyacetate and has a cubic crystalline structure. Normal zinc-acetate crystallizes in cubic volume centered lattice with a parameter $a = 10.92 \pm 0.05$. The properties of normal zinc acetate differ substantially from those of the normal beryllium acetate. The analogy between zinc- and beryllium acetate-compounds comprises only their oxy-acetates; whereas the normal acetates differ considerably. There are 1 table and 5 references, 3 of which are Soviet.

Card 1/2

On the Acetates of Zinc and Beryllium

78-3-6-28/30

SUBMITTED: November 21, 1977

AVAILABLE: Library of Congress

1. Zinc acetate--Synthesis
2. Beryllium acetate--Synthesis

Card 2/2

AUTHORS: Maksimov, V. N., Semenenko, K. N. 78-3-6-29/30

TITLE: On Lanthanum-Acetate With 5 mol Water (O pyativodnom atsetate lantana)

PERIODICAL: Zhurnal Neorganicheskoy Khimii, 1958, Vol. 3, Nr 6, pp. 1468-1469 (USSR)

ABSTRACT: A crystalline lanthanum acetate with 5 mol water was obtained by slow crystallization at temperatures of 13 to 20°C. The crystals are big and prismatic. The chemical composition of this compound is the following: (in per cent by weight) La_2O_3 - 40,3, C - 17,85, H - 5,00. The parameters of the elementary cells were found by radiographic investigations: $a = b = 9,0 \pm 0,1 \text{ kX}$, $c = 8,9 \pm 0,1 \text{ kX}$, $\alpha = \beta = \gamma = 96 \pm 1^\circ$. The thermographic analyses of the compound were investigated and it was found that the compound loses 17,7 % of its weight at 25°C and passes over into lanthanum acetate with 1 mol water. The last mol water is delivered at 150°C. The anhydrous lanthanum acetate is stable up to 250°C and beyond this temperature the decomposition continues white $\text{LaO} \cdot \text{CH}_3 \text{COO}$

Card 1/2

On Lanthanum-Acetate With 5 mol Water

78-3-6-29/30

forms. There are 1 figure and 4 references, 1 of which is Soviet.

SUBMITTED: November 22, 1957

AVAILABLE: Library of Congress

1. Chemical compound--Properties
2. Chemical compound--Analysis
3. Lanthanum acetate--Applications

Card 2/2

AUTHORS: Zlomanov, V.P., Novoselova, A.V., SOV/78-3-7-1/44
Pashinkin, A.S., Simanov, Yu.P., Semenerko, K.N.

TITLE: Determination of the Pressure of Steam Saturated With Solid
Tellurium Dioxide (Opređoleniye davleniya nasyshchennogo para
tverdogo dvuokisi tellura)

PERIODICAL: Zhurnal neorganicheskoy khimii, 1958, Vol. 3, Nr 7, pp 1473-1477
(USSR)

ABSTRACT: The pressure of steam saturated with solid tellurium dioxide was
determined in the temperature interval of 457-704° C by means of
a radioactive tellurium isotope. The phase composition of telluri-
um dioxide was determined, for which purpose thermograms for the
temperature interval of 25-800° C, as well as heating- and cool-
ing diagrams were made. X-ray analyses showed that the crystal
lattice of tellurium dioxide is tetragonal and has the follow-
ing parameters: $a = 4.756$, $c = 7.588$ kX.
On the strength of the results obtained by thermographical and
radiographical analyses it follows that the solid phase of the
vaporous tellurium dioxide shows tetragonal modifications. There
are 3 figures, 2 tables, and 16 references, 9 of which are Soviet.

Card 1/2

Determination of the Pressure of Steam Saturated With
Solid Tellurium Dioxide

SOV/ 78-3-7-1/44

ASSOCIATION: Moskovskiy gosudarstvennyy universitet im. M.V.Lomonosova
(Moscow State University imeni M.V.Lomonosov)

SUBMITTED: July 8, 1957

1. Steam--Pressure 2. Pressure--Determination 3. Tellurium
dioxide--Phase studies 4. Tellurium isotopes--Applications
5. X-rays--Applications

Card 2/2

AUTHORS: Grigor'yev, A.I., Novoselova, A.V., Semenko K.N. SN/78-3-7-22/44

TITLE: On the Compounds of Berylliumoxyacetate With Ethylamine and Butylamine (O sovedinerdyakh oksiatsetata beriliya s etilaminom i butilaminom)

PERIODICAL: Zhurnal neorganicheskoy khimii, 1958, Vol. 3, Nr 7, pp 1599-1604 (USSR)

ABSTRACT: Compounds of berylliumoxyacetate with ethylamine and butylamine were synthesized. Analyses resulted in the following compositions:
 $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_{6.8} \text{C}_2\text{H}_5\text{NH}_2$, $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_{6.4} \text{C}_2\text{H}_5\text{NH}_2$,
 $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_{6.3} \text{C}_2\text{H}_5\text{NH}_2$, $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_{6.8} \text{C}_4\text{H}_9\text{NH}_2$,
 $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_{6.4} \text{C}_4\text{H}_9\text{NH}_2$, $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_6 3 \text{C}_4\text{H}_9\text{NH}_2$ and
 $\text{Be}_4(\text{CH}_3\text{COO})_6 \text{C}_4\text{H}_9\text{NH}_2$.
 The complex compounds with butylamine are easily decomposed. The crystal lattices of the compounds $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_{6.4} \text{C}_2\text{H}_5\text{NH}_2$ and $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_{6.4} \text{C}_4\text{H}_9\text{NH}_2$ are very similar. The thermograms of the compounds of berylliumoxyacetate with ethylamine and butylamine

Card 1/2

AUTHORS: Novoselova, A. V., Semenenko, K. N. SOV/78-3-9-34/38

TITLE: Chlorination of Beryllium Oxide With Carbon Tetrachloride
(Khlorigirovaniye okisi berilliya chetyrekhlolistym uglerodom)

PERIODICAL: Zhurnal neorganicheskoy khimii, 1958, Vol 3, Nr 9, pp 2213-2214
(USSR)

ABSTRACT: An apparatus for the chlorination of beryllium oxide with carbon tetrachloride in the so-called "boiling layer" was described. Beryllium oxide is chlorinated at a temperature of 700-800°C in a quartz boat with carbon tetrachloride vapor and with nitrogen as a carrier. The beryllium chloride formed - BeCl_2 - is removed from the zone of reaction by sublimation. The reactor for the reaction is placed vertically. This apparatus for chlorination of beryllium oxide in the boiling layer shows that after 20 to 25 minutes almost 95% of all the beryllium oxide has been chlorinated. A chlorination of beryllium oxide in a horizontal reactor, however, does not supply a quantitative yield even after several days. There are 1 figure and 3 references, 0 of which is Soviet.

Card 1/2

AUTHORS: Grigor'yev, A. I., Semenenko, K. N. SOV/78-3-12-34/36

TITLE: Concerning the Compound Beryllium Oxyacetate With Methyl Amine
(O soyedinenii oksiatsetata berilliya s metilaminom)

PERIODICAL: Zhurnal neorganicheskoy khimii, 1958, Vol 3, Nr 12,
pp 2806-2807 (USSR)

ABSTRACT: The compound $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_6 \cdot 4\text{CH}_3\text{NH}_2$ was prepared in the form of large rhombic prisms by slow vaporization of a saturated solution in liquid methyl amine at room temperature. X-ray studies showed that this compound crystallizes with a triclinic lattice with the following parameters: $a=8.34 \text{ \AA}$, $b=10.20 \text{ \AA}$, $c=9.44 \text{ \AA}$, $\alpha=58^\circ$, $\beta=55^\circ$, $\gamma=60^\circ$. The polythermal decomposition curve of $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_6 \cdot 4\text{CH}_3\text{NH}_2$ was constructed using the thermogravimetric determination method. The compounds formed by beryllium with ethyl and butyl amines have similar properties to those which form from methyl amine. There are 1 figure and 2 Soviet references.

Card 1/2

SOV/156-58-4-22/49

AUTHORS: Tarasevich, V. I., Semenenko, K. A., Semenenko, K. N.

TITLE: The Radiographic Investigation of the Products of Chemical Reactions in Spectroscopic Determinations of Niobium
(Rentgenograficheskoye izucheniye produktov khimicheskikh reaktsiy pri spektral'nom opredelenii niobiya)

PERIODICAL: Nauchnyye doklady vysshey shkoly. Khimiya i khimicheskaya tekhnologiya, 1958, Nr 4, pp 700-705 (USSR)

ABSTRACT: In the present paper the products formed on the carbon electrode in the spectrum analysis of niobium determination were radiographically investigated. The following samples were investigated: I. Nb+C; II. Si+C; III. Nb+Si+C; IV. Nb₂O₅+C; V. Nb₂O₅+Si+C; VI. Nb₂O₅+SiO₂+C. The radiograms of the products were taken. The radiographic investigation shows that in the interaction of metallic niobium with carbon only niobium carbide is formed. In the interaction between silicon and carbon SiC is formed. The interaction between carbon and a mixture of Nb and Si takes a complex way, however. The radiographic analysis shows that in the reaction products the

Card 1/3

SOV/156-58-4-22/49

The Radiographic Investigation of the Products of Chemical Reactions in
Spectroscopic Determinations of Niobium

following phases are formed: cubic NbC and tetragonal β -Nb₅Si₃. The interaction of Nb₂O₅ with the carbon electrode shows only a modification of Nb₂O₅ with small impurities of NbO₂. Lower niobium oxides were not determined. In the interaction between niobium pentoxide Nb₂O₅ and elementary silicon with carbon NbO₂ and a phase difficultly identified are formed. The interaction between niobium pentoxide Nb₂O₅ and silicon dioxide with carbon leads to the formation of NbO₂ and niobium pentoxide. In the presence of elementary silicon and SiO₂ niobium dioxide is formed on the carbon crater during the spectroscopic determination of niobium. The excitation source and excitation conditions as well as the amperage do practically not exert any influence upon the composition of niobium phases in the carbon crater. There are 1 figure, 4 tables, and 6 references, 4 of which are Soviet.

Card 2/3

5(2)

AUTHOR: Semenenko, K.N.

SOV/55-58-5-24/34

TITLE: On Oxysalts of Beryllium (Ob oksisolyakh berilliya)

PERIODICAL: Vestnik Moskovskogo universiteta, Seriya matematiki, mekhaniki, astronomii, fiziki, khimii, 1958, Nr 5, pp 151 - 164 (USSR)

ABSTRACT: The present paper contains a short survey of the methods of production and of the properties of the so-called oxysalts $\text{Be}_4\text{O}(\text{RCOO})_6$, as well as of the applicabilities of these combinations. It mainly concerns with a summary of western results. The following Soviet researchers were mentioned: A.V. Novoselova, A.I. Grigor'yev, K.N. Semenenko, V.N. Maksimov, N.Ya. Turova, and N.N. Krasovskaya. The paper contains no new results. There are 55 references, 12 of which are Soviet, 13 French, 12 German, 9 English, 6 American, 2 Japanese, and 1 Italian.

ASSOCIATION: Kafedra neorganicheskoy khimii (Chair of Inorganic Chemistry)

SUBMITTED: November 4, 1957

Card 1/1

5 (2, 3)

AUTHORS:

Novoselova, A. V., Semenenko, K. N.,
Turova, N. Ya.

SOV/55-58-6-18/31

TITLE:

Beryllium-organic Compounds (Berilliyorganicheskiye
soyedineniya)

PERIODICAL:

Vestnik Moskovskogo universiteta. Seriya matematiki,
mekhaniki, astronomii, fiziki, khimii, 1958, Nr 6,
pp 139-147 (USSR)

ABSTRACT:

This article gives a survey of the known possibilities of obtaining and of the fundamental properties of beryllium-organic compounds. The data concerning these compounds are from Western publications. The various existing types of beryllium-organic compounds (BeR_2 and BeRX , where R denotes aliphatic or aromatic radicals, X - a halide, alkoxyl, hydrogen or the NR_2 group), as well as the hitherto nearly unknown type RBeR' are discussed in detail. There are 2 tables and 40 references, 4 of which are Soviet.

Card 1/2

Beryllium-organic Compounds

SOV/55-58-6-18/31

ASSOCIATION: Kafedra neorganicheskoy khimii (Chair for Inorganic Chemistry)

SUBMITTED: December 25, 1957

Card 2/2

Semenenko K. N.

AUTHORS: Simanov, Yu.F., Burov, V.S., (deceased),
Semenenko, K.N.

32-43/55

TITLE: A Simple Device for Taking X-Ray Pictures of Monocrystals at
High Temperatures (Prostoye prispobleniye dlya rentgenos'yemki
monokristalla pri vysokoy temperature).

PERIODICAL: Zavodskaya Laboratoriya, 1958, Vol. 24, Nr 1, pp. 107-107 (USSR)

ABSTRACT: The present paper describes a new additional device to be used
with the X-ray camera PKCO. According to the diagram the device
consists of a round copper block with an opening passing through
its center, into which a thermocouple is fitted. This block rests
upon a second asbestos block which, at the same time, serves as a
current- and heat insulator. The device is placed firmly upon the
goniometer head of the camera. The ends of the thermocouple are
connected to the galvanometer "M-82". On the copper block an annu-
lar Nichrome heater is fastened which is connected with the mains
by means of a step-down transformer. The sample (monocrystal) is
pasted immediately on to the soldering seams of the thermocouple
by means of a special adhesive, which should be somewhat above

Card 1/2

A Simple Device for Taking X-Ray Pictures of
Monocrystals at High Temperatures

32-1-43/55

the openings of the copper block. In this way the monocrystal of $\text{Be}_4\text{O}(\text{GH}_3\text{GOO})_6$ was investigated at temperatures of 20, 50, 80 and 120°. The transformation of crystallization could be well determined by means of X-ray pictures. There are 1 figure, and 2 references, 1 of which is Slavic.

AVAILABLE: Library of Congress

Card 2/2 1. X-ray cameras-Adapters

SEMINOV, K. N.

AUTHORS: Grigor'ev, A. I., Novoselova, A. V.,
Semenenko, K. N.

22-2-25/60

TITLE: Determination of the Molecular Weight of Dissolved Substances According to the Method of Diffusion Through a Porous Glass Platelet (Opredeleniye molekulyarnykh veshchestv metodom diffusii cherez poristuyu steklyannuyu plastinku)

PERIODICAL: Zavodskaya Laboratoriya, 1958, Vol. 24, Nr 2, pp. 190-192 (USSR)

ABSTRACT: The idea of Northrop Anson (reference 1) was applied to determine molecular weights (of the order of magnitude of 400 to 500) of substances dissolved in chloroform. The molecular weights are computed from the experimentally found diffusion coefficients of the substance under investigation and of a substance with a known molecular weight with the formula:

$$\frac{D'}{D''} = \frac{\sqrt{M''}}{\sqrt{M'}}$$

Card 1/2

The oxycetate and the oxypropionate of beryllium were in-

Determination of the Molecular Weight of Dissolved Substances 32-2-25/60
According to the Method of Diffusion Through a Porous Glass Platelet

investigated and a difference of only 0.5% to the computed molecular weight was found. When the molecular weight of anthracene was determined, however, a difference of 19% was found, which can be due to the differences between the structures of the Be-oxyacetate and that of anthracene (corresponding to the observations made by Drintzinger, reference 3). The application of a standard as an accompanying substance is therefore proposed for the purpose of improving the method. The radioactive C^{14} isotope was, among others, used in order to remove difficulties of analytical kind. For the determination of the specific activities, the solutions within the cell and without were vaporized after diffusion, the residue was dessiccated, combusted and the $C^{14}O_2$ was transformed into $BaC^{14}O_3$. There are 1 figure, 4 tables, and 4 references, 1 of which is Slavic.

ASSOCIATION: Moscow State University imeni M. V. Lomonosov
(Moskovskiy gosudarstvennyy universitet imeni M. V. Lomonosova)

AVAILABLE: Library of Congress
Card 2/2 1. Molecular weight-determination 2. Chloroform-Applications

SOV/20-122-3-20/57

AUTHORS: Grigor'yev, A. I., Corresponding Member, Academy of Sciences,
USSR, Novoselova, A. V., Semenenko, K. N.

TITLE: Compounds of Beryllium Hydroxy Acetate With Sulfur Dioxide
(Soyedineniya oksiatsetata berilliya s sernistym angidridom)

PERIODICAL: Doklady Akademii nauk SSSR, 1958, Vol 122, Nr 3, pp 397-399
(USSR)

ABSTRACT: Affiliation products containing mainly amino nitrogen are de-
scribed for beryllium hydroxy acetate (Refs 1-3). These com-
pounds are stable enough and are probably formed at the ex-
pense of the free electron pair of nitrogen. However, for the
substance mentioned first in the title compounds of a weaker
binding may be expected, namely of the type of the so-called
"inclusion compounds" (soyedineniya vklyucheniya Pl.) (Refs
4,5). The compound mentioned in the title probably is such an
"inclusion compound" (Ref 2). This problem is discussed in
detail in the present paper. In the concentration by the
evaporation of a solution of beryllium hydroxy acetate in
liquid sulfur dioxide the latter compound is precipitated in
form of well developed octahedrons. The thus forming compound

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Compounds of Beryllium Hydroxy Acetate With Sulfur Dioxide

SOV/20-122-3-20/57

is extremely instable at room temperature and decomposes into its two initial products. This makes difficult the determination of its composition and the preparative isolation by means of the usual methods of chemical analysis. In order to investigate the interaction of both substances mentioned in the title the authors studied the method of the construction of diagrams at a constant temperature: composition - vapor pressure in the system formed by them. For this purpose they used the Huettig tensiometer (тензиевдиометр) (Ref 6). The working process is described. The equilibrium could be observed after 10-20 hours. Figure 1 shows the isothermal lines of the composition versus pressure function for -9,5, -15, -20 and -30°. From the general view of the isothermal lines it can be seen that in the case of a concentration by evaporation of one of the mentioned saturated solutions a compound $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_6 \cdot 2\text{SO}_2$ is precipitated. Thus, it was observed that the compound $3\text{Be}_4\text{O}(\text{CH}_3\text{COO})_6 \cdot 4\text{SO}_2$ (described by the authors in reference 2) represents a product of a partial decomposition of the compound of beryllium hydroxy acetate with molecules of sulfur dioxide. Besides a compound 2 : 1 another one 1 : 1 $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_6 \cdot \text{SO}_2$ which forms on the

Card 2/4

SOV/20-122-3-20/57

Compounds of Beryllium Hydroxy Acetate With Sulfur Dioxide

decomposition of the first was observed in the system. The stability of the compounds decreases with the increase in temperature. It is remarkable that in this case the values of the heats of formation (on the average 9,22 kcal per 1 g mol SO_2) are lower than in the normal case of coordination compounds. Furthermore, it is of interest that at -10° (boiling point of SO_2) the discussed compounds have the characteristic features of solid solutions and can exist only at increased pressure. According to radiographic analyses $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_6 \cdot 2\text{SO}_2$ crystallizes in a cubic diamond-like lattice with a period of the elementary cell of $a = 17,1 \text{ \AA}$. The density at $-12^\circ = 1,43$, roentgen density = 1,42. In conclusion a rough outline of the structure of this substance is given. There are 2 figures and 7 references, 4 of which are Soviet.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet im. M. V. Lomonosova
(Moscow State University imeni M. V. Lomonosov)

Card 3/4

5(2)

AUTHORS:

Turova, N. Ya., Novoselova, A. V., Semenenko, K. N.,
Savost'yanova, R. I.

SOV/78-4-3-10/34

TITLE:

On the Phenolates of Beryllium (O fenolyatakh berilliya)

PERIODICAL:

Zhurnal neorganicheskoy khimii, 1959, Vol 4, Nr 3,
pp 549-552 (USSR)

ABSTRACT:

The interaction between beryllium chloride and β -naphthol and p- and m-cresols has been investigated and the properties of the resulting phenolates have been described. The reaction of beryllium chloride with o-, p-, m-cresol takes place at 90-100°C. The interaction of p- and m-cresol with BeCl_2 takes place under a strong development of HCl. The phenolates of beryllium are white, amorphous, hygroscopic substances, which slowly hydrolyze in air. Thermographic and radiographic investigations were carried out. The following phenolates have been prepared: β -naphthol beryllium ($\text{Be}(\text{OC}_{10}\text{H}_7)_2$) and $\text{Be}(\text{p-OC}_7\text{H}_7)_2$ and $\text{Be}(\text{m-OC}_7\text{H}_7)_2$. The phenolates of beryllium are slightly soluble in benzene and xylene, stable in ether. Decomposition occurs under the action of methyl alcohol.

Card 1/2

SOV/78-4-3-10/34

' On the Phenolates of Beryllium

There are 1 figure and 7 references, 2 of which are Soviet.

SUBMITTED: January 4, 1958

Card 2/2

5(2)

SOV/78-4-4-41/44

AUTHORS:

Semenenko, K. N., Gordeyev, I. V.

TITLE:

Investigation of the Monoclinic Modification of Beryllium Oxyacetate (Issledovaniye monoklinnoy modifikatsii oksiatsetata berilliya)

PERIODICAL:

Zhurnal neorganicheskoy khimii, 1959, Vol 4, Nr 4, pp 952-954 (USSR)

ABSTRACT:

The monoclinic modification of beryllium oxyacetate is transformed very slowly into the stable cubic modification, whereby the vapor pressure may be determined according to Knudsen's method. The vapor pressure of the monoclinic modification dependent on temperature is expressed by the equation

$$\log P = 12.777 - \frac{6025.2}{T}$$

The results of measurement of the vapor pressure are contained in table 1. The reciprocal position of the straight line

$\lg P = a - \frac{B}{T}$ of the three modifications of $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_6$ is given in figure 2. The heat of sublimation of the monoclinic modification $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_6$ amounts to 27.56 kcal/mole. The heats

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Investigation of the Monoclinic Modification of Beryllium Oxyacetate SOV/78-4-4-41/44

of sublimation of the monoclinic modification and high-temperature modification of beryllium oxyacetate are close to one another - 27.56 and 27.10 kcal/mole. They indicate great structural similarity of both modifications. The authors thank A. S. Pashinkin for valuable advice and assistance in the work. There are 2 figures, 1 table, and 6 references, 2 of which are Soviet.

SUBMITTED: October 9, 1958

Card 2/2

5(2)

AUTHORS: Turova, N. Ya., Novoselova, A. V., SOV/78-4-5-10/46
Semenenko, K. N.

TITLE: On the Alcoholates of Beryllium (Ob alkogolyatakh berilliya)
Communication I.

PERIODICAL: Zhurnal neorganicheskoy khimii, 1959, Vol 4, Nr 5,
pp 997-1001 (USSR)

ABSTRACT: The syntheses for the preparation of the alcoholates of beryllium were investigated and some properties of beryllium ethylate were discussed. The reaction between metallic beryllium and absolute ethyl alcohol was recommended in the presence of BeCl_2 , HgCl_2 or J for the purpose of synthesizing beryllium ethylate. Beryllium ethylate of the composition $\text{Be}(\text{OC}_2\text{H}_5)_2$ is a white amorphous substance. The product is not soluble in water and in the usual organic solvents. Several properties of beryllium ethylate, especially its behavior with respect to absolute ethyl alcohol, anhydrous acetic acid, and alcoholic BeCl_2 -solution were investigated. In the interaction between beryllium ethylate and anhydrous acetic acid in an ether medium an insoluble compound with the composition $\text{Be}(\text{OC}_2\text{H}_5)(\text{OCOCH}_3)$ is formed after some hours with a molar ratio of components of 1:1. In the interaction of beryllium

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On the Alcoholates of Beryllium

SOV/78-4-5-10/46

ethylate with 0.5 N beryllium chloride solution, beryllium ethylate dissolves completely in absolute ethyl alcohol within two hours. With the evaporation of this solution a syrup-like mass is formed. The dissolution process of beryllium ethylate is connected with the formation of complex compounds of the type $\text{Be}(\text{OR})_2 \cdot \text{BeCl}_2$ or $\text{Be}(\text{BeCl}_2(\text{OR})_2)$ in alcoholic BeCl_2 -solution. The interaction between beryllium and absolute methyl alcohol shows that, in the presence of BeCl_2 , HgCl_2 and J, a compound of variable composition is formed, for which the general formula $[\text{xBe}(\text{OCH}_3)_2 \cdot \text{yBe}(\text{OCH}_3)\text{Hal}]_n$ holds. In the interaction between BeCl_2 and $\text{Na}[\text{Be}(\text{OCH}_3)_4]$ a mixture of NaCl and $\text{Be}(\text{OCH}_3)_2$ is formed. There are 13 references, 3 of which are Soviet.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet im. M. V. Lomonosova
(Moscow State University imeni M. V. Lomonosov)

SUBMITTED: February 3, 1958

Card 2/2

5(2)

AUTHORS:

Turova, N. Ya., Novoselova, A. V., SOV/78-4-5-45/46
Semenenko, K. H.

TITLE:

The Synthesis of the Etherates of Beryllium Halides (Sintez
efiratorov galogenidom berilliya)

PERIODICAL:

Zhurnal neorganicheskoy khimii, 1959, Vol 4, Nr 5,
pp 1215-1216 (USSR)

ABSTRACT:

The synthesis of the etherates of beryllium chloride and beryllium bromide is carried out by the direct interaction between the metallic beryllium and halogens or hydrogen chloride in an absolute ether medium. The syntheses of the etherates of bromides and iodides of beryllium are carried out in the nitrogen flow. The following compounds were isolated: $\text{BeCl}_2 \cdot 2(\text{C}_2\text{H}_5)_2\text{O}$ and $\text{BeBr}_2 \cdot 2(\text{C}_2\text{H}_5)_2\text{O}$. The melting temperatures of the etherates of beryllium chloride and beryllium bromide agree well with the data given by the authors for preparations produced by the interaction of anhydrous halides with ether. It was not possible to represent etherate of beryllium iodide in the purest form. The method described for the synthesis of the etherates of beryllium halides is of a general character and may be used for the production of

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5(2)

SOV/78-4-10-38/40

AUTHORS: Turova, N. Ya., Novoselova, A. V., Semenenko, K. N.

TITLE: On Compounds of Beryllium Chloride With Tetrahydrofuran

PERIODICAL: Zhurnal neorganicheskoy khimii, 1959, Vol 4, Nr 10,
pp 2410 - 2411(USSR)

ABSTRACT: The system BeCl_2 - tetrahydrofuran was investigated in the temperature range - 78 up to +150° (Table 1: Solubility, Fig 1: Phase Diagram, Fig 2: Dependence of logC on $\frac{1}{T}$). At low temperature $\text{BeCl}_2 \cdot 3\text{C}_4\text{H}_8\text{O}$ is formed as solid phase at the bottom which decomposes at -2° to yield $\text{BeCl}_2 \cdot 2\text{C}_4\text{H}_8\text{O}$. This melts at 150° without decomposition, is more stable than the etherate of beryllium chloride when exposed to air, well soluble in benzene and insoluble in petroleum ether. There are 2 figures, 1 table, and 4 references, 3 of which are Soviet.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet im. M. V. Lomonosova
(Moscow State University imeni M. V. Lomonosov)

SUBMITTED: May 22, 1959
Card 1/1

5(2)

SOV/20-125-3-25/63

AUTHORS:

Grigor'yev, A. I., Novoselova, A. V., Corresponding Member
AS USSR, Semenenko, K. N.

TITLE:

On the Compound Formed by Beryllium Oxy-acetate and Nitrogen
Dioxide (O soyedinenii oksiatsetata berilliya s dvuokis'yu
azota)

PERIODICAL:

Doklady Akademii nauk SSSR, 1959, Vol 125, Nr 3, pp 557-559
(USSR)

ABSTRACT:

It was found that $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_6$ forms adducts with NO_2 , similar
to those formed with SO_2 ($\text{Be}_4\text{O}(\text{CH}_3\text{COO})_6 \cdot 2\text{SO}_2$ and
 $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_6 \cdot \text{SO}_2$). The mentioned oxy-acetate is well soluble
in liquid nitrogen dioxide at room temperature. If this solu-
tion is vaporized achromatic needlelike anisotropic crystals are
separated. They decompose quickly in air under formation of
brown NO_2 -vapors. After this decomposition beryllium oxy-
acetate is left back in its cubical basic modification. The
composition of the crystals may be approximately described by
the following formulas: $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_6 \cdot 3\text{NO}_2$ or

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SOV/20-125-3-25/63

On the Compound Formed by Beryllium Oxy-acetate and Nitrogen Dioxide

$\text{Be}_4\text{O}(\text{CH}_3\text{COO})_{6.1,5}\text{N}_2\text{O}_4$. By means of the measurement of the magnetic susceptibility was found that N_2O_4 probably takes part in the mentioned compounds. In order to define their composition precisely as well as in order to determine the possibility of formation of other compounds in the system beryllium oxy-acetate - nitrogen dioxide the diagrams: composition - vapor pressure at constant temperature were plotted. The tensi-eudiometer of Hüttig (Khyuttig) served for this purpose. Its main parameters and the method were the same, as in reference 1 with the exception of a small modification which takes the aggressiveness of the gas into account because it reacts with mercury. After 2-3 hours the equilibrium in the system was re-established. Figure 1 shows isothermal lines at 10.0 and 19.0°C. Their general shape shows that the compound $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_{6.1,5}\text{N}_2\text{O}_4$ is separated by the evaporation of the saturated $\text{Be}_4\text{O}(\text{CH}_3\text{COO})_6$ solution in the liquid NO_2 . No other compounds were found to exist in the system. The last mentioned compound dis-

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SOV/20-125-3-25/63

On the Compound Formed by Beryllium Oxy-acetate and Nitrogen Dioxide

sociates as a true chemical compound in contrast to the two compounds formed with SO_2 (mentioned above). The decomposition of the two last mentioned compounds proceeds by the formation of phases of variable composition. This contrast is assumed to explain the quicker establishment of the dissociation equilibrium of the compound with NO_2 . The dependence of the dissociation pressure was explained by the isothermal lines (Table 1, Fig 2). Furthermore the compound obtained was investigated radiographically, its density and crystalline structure determined. There are 2 figures, 1 table, and 4 references, 1 of which is Soviet.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet im. M. V. Lomonosova
(Moscow State University imeni M. V. Lomonosov)

SUBMITTED: January 2, 1959

Card 3/3

SESTANO, R.N.

High-temperature modification of beryllium acetate. Zhur.
strukht. khim. 1 no. 4:42-44 1960. (NII 14:2)

1. Moskovskiy gosudarstvennyy universitet imeni M.V. Lomonosova.
(Beryllium acetate)

TUROVA, N.Ya.; NOVOSELOVA, A.V.; SEMENENKO, K.W.

Solubility in the system beryllium chloride dietherate-
ethyl ether. Zhur.neorg.khim. 5 no.1:117-123 Ja '60.
(MIRA 13:5)

1. Moskovskiy gosudarstvennyy universitet im. M.V.Lomonosova.
(Beryllium compounds) (Ether)

SEMENENKO, K.N.; TUROVA, N.Ya.; URAZBAYEVA, R.N.

Synthesis of the lithium aluminum hydride. Zhur.neorg.khim.
5 no.2:508 F '60. (MIRA 13:6)

1. Moskovskiy gosudarstvennyy universitet im. M.V.Lomonosova.
(Aluminum lithium hydride)

69047

S/078/60/005/03/009/048
B004/B002

5.3700(B)
AUTHORS: Maksimov, V. N., Semenenko, K. N.,
Naumova, T. N., Novoselova, A. V.

TITLE: Aluminum Acetates¹

PERIODICAL: Zhurnal neorganicheskoy khimii, 1960, Vol 5, Nr 3, pp 558 - 564
(USSR)

ABSTRACT: After a brief survey of publications, the authors report on their investigation of aluminum acetates. They produced aluminumtriacetate from aluminum ethylate and acetic anhydride. $\text{Al}(\text{CH}_3\text{COO})_3$ is easily soluble in liquid ammonia under the development of $\text{Al}(\text{CH}_3\text{COO})_3 \cdot 3\text{NH}_3$.

During thermal decomposition, the triacetate gradually passes over into di- and monoacetate (Figs 1,2). The data of the radioanalysis taken by means of an RKD camera and Fe radiation of the BSV tube are given by table 2. The authors also investigated basic aluminum acetates. From $\text{Al}(\text{OH})_3$ plus acetic acid and also from AlCl_3 plus acetic acid they obtained the same compound $\text{Al}(\text{OH})(\text{CH}_3\text{COO})_2$ whose radioanalysis is given in table 1. The basic diacetate has a rhombic, face-centred lattice with the lattice constants being $a = 13.62 \pm 0.01 \text{ \AA}$, $b = 14.40 \pm 0.01 \text{ \AA}$, $c = 12.60 \pm 0.01 \text{ \AA}$. On the basis

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Aluminum Acetates

69047

S/078/60/005/03/009/048
B004/B002

of the density being 1.67, a lattice cell contains 16 molecules. The basic diacetate is little soluble in water, chloroform and liquid SO_2 , and insoluble in alcohol, acetone, ether, and liquid ammonia. On the basis of the thermogram (Fig 3) taken by means of the Kurnakov pyrometer type PK-42, the formula $\text{Al}(\text{OH})(\text{CH}_3\text{COO})_2$ was found to be right, not $\text{Al}_2\text{O}(\text{CH}_3\text{COO})_4 \cdot \text{H}_2\text{O}$. During the reaction of sodium acetate (or barium acetate) and aqueous solutions of AlCl_3 , a basic salt was obtained whose composition is between $\text{Al}(\text{OH})(\text{CH}_3\text{COO})_2 \cdot 2\text{H}_2\text{O}$ and $\text{Al}(\text{OH})(\text{CH}_3\text{COO})_2 \cdot 2.5\text{H}_2\text{O}$, and whose radiogram (Table 2) differs from that of $\text{Al}(\text{OH})(\text{CH}_3\text{COO})_2$. The thermogram of figure 4 shows the water separation of this salt during heating. The nonaqueous salt thus developing, however, radiographically differs from the salt produced by means of free acetic acid, despite the same stoichiometric composition. By the influence of sodium acetate on aluminum sulphate, the compound $\text{Al}(\text{OH})(\text{CH}_3\text{COO})_2 \cdot 2.5\text{H}_2\text{O}$ was obtained, and during the reaction of sodium acetate and aluminum nitrate, $\text{Al}(\text{OH})(\text{CH}_3\text{COO})_2$ developed; both were radiographically identified. Aluminum nitrate with acetic anhydride developed a

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Aluminum Acetates

69047

S/078/60/005/03/009/048
B004/B002

compound of varying composition which always contained up to 3% NO_3 , and whose radiogram was identical with that of aluminum triacetate. There are 2 figures, 4 tables, and 22 references, 4 of which are Soviet. 4

SUBMITTED: November 22, 1958

Card 3/3

TUROVA, N.Ya., NOVOSELOVA, A.V., SEMENENKO, K.N.

Solubility in the system beryllium bromide dietherate - ether.
Zhur. neorg. khim. 5 no.4:941-944 Ap '60. (MIRA 13:7)

1. Moskovskiy gosudarstvennyy universitet im. M. V. Lomonosova.
(Beryllium compounds)

SEMENENKO, K.N.

Structure of the inclusion compound of beryllium oxyacetate
with sulfur dioxide. Zhur. neorg. khim. 5 no. 12:2687-2689
D '60. (MIRA 13:12)

(Beryllium compounds)

86488

15.8117

2209, 2109, 2808, 3215

S/078/60/005/008/021/031/XX
B023/B066

AUTHORS: Turova, N. Ya.; Novoselova, A. V., Semenenko, K. N.

TITLE: Compounds of Beryllium Chloride With Ethers

PERIODICAL: Zhurnal neorganicheskoy khimii, 1960, Vol. 5, No. 8,
pp. 1705-1709

TEXT: The authors report on their synthesis and investigation of new compounds of beryllium chloride with dimethyl ether, dibutyl ether, tetrahydropyran, and ethylene glycol dimethyl ether (1,2-dimethoxy-ethane). The following rules were established: 1) The thermal stability of beryllium chloride complexes with ethers of monovalent radicals increases rapidly on transition of compounds of the aliphatic series to cyclic ethers. 2) The melting point of $\text{BeCl}_2 \cdot 2\text{R}_2\text{O}$ (R = alkyl radical) decreases considerably in the homologous series of aliphatic ethers (at R = CH_3 , C_2H_5 , $n\text{-C}_4\text{H}_9$, the melting point is 63° , 43° , and $<-70^\circ\text{C}$, respectively). 3) The largest difference may be seen in beryllium chloride compounds with ethers

Card 1/3

Compounds of Beryllium Chloride With Ethers

36488

S/078/60/005/008/021/031/XX
E023/B066

of mono- and divalent alcohols. Complexes with monocethers form well crystallizable compounds which melt congruently (except $\text{BeCl}_2 \cdot 2(\text{CH}_3)_2\text{O}$) and are soluble in the corresponding ethers, in aromatic hydrocarbons, carbon tetrachloride, and carbon disulfide, but insoluble in paraffin hydrocarbons. On the other hand, compounds of the beryllium halides with ethers of divalent alcohols are decomposed on heating without melting. They are insoluble in all organic solvents, except in alcohols, and are inert to a higher degree than compounds with ethers of monovalent alcohols. All these properties indicate that compounds of beryllium chloride with ethers of divalent alcohols have a polymeric character. Beryllium chloride dioxanate is assumed to have a chain structure formed by the coordination bonds of both oxygen atoms in dioxane. Well soluble and crystallizable monodioxanates, such as chloride, bromide, and iodide dioxanates of divalent mercury are, apparently, polymers with a low association degree. The formation of didioxanates is particularly characteristic of halides of divalent metals having the coordination number 6. The properties of magnesium, nickel, and iron halides, as well as of calcium, strontium, and barium bromides and iodides are similar to those of monodioxanates.

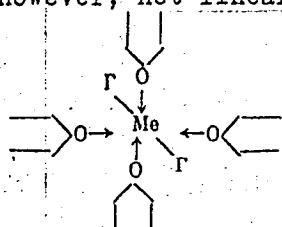
Card 2/3

86488

Compounds of Beryllium Chloride With Ethers

S/078/60/005/008/021/031/XX
E023/B066

They are insoluble in dioxane and organic solvents, and separate in the form of a fine crystalline powder. They also have a polymeric structure which is, however, not linear but reticular:



ZNC 1960, Vol. 5, No. 8,
pp. 1705-1709

X = Halogen. There are 4 tables and 14 references: 8 Soviet, 1 US, and 5 German.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet im. M.V. Lomonosova
(Moscow State University imeni M. V. Lomonosov)

SUBMITTED: May 22, 1959

Card 3/3

S/189/60/000/003/008/013/XX
B033/B067

AUTHORS: Tarasevich, N. I., Semenenko, K. A., Semenenko, K. N.
TITLE: X-Ray Photographic Method of Determining the Products of
Chemical Reactions in the Spectral Determination of Tantalum ✓
PERIODICAL: Vestnik Moskovskogo universiteta. Seriya 2, khimiya, 1960, 15,
No. 3, pp. 37-39

TEXT: The authors studied the reaction products which were formed from tantalum pentoxide in the electric arc in the crater of the carbon electrode (Ta_2O_5). The investigation method applied is described in an earlier paper. (Ref. 1). The very finely powdered Ta_2O_5 was filled into the electrode crater and closed with a cover of coal (provided with an opening for the gases). In all experiments the reaction conditions in the arc were the same. The X-ray powder patterns of the reaction products were taken with PKD (RKD) cameras. A ECB (BSV) tube served as radiation source (copper electrode). The product formed from Ta_2O_5 (under the

Part 1/3

SEMENENKO, K.N.; KURDYUMOV, G.M.

Some properties of beryllium α -hydroxynaphthoate. Vest. Mosk. un.
Ser. 2: Khim. 15 no.5:56-58 S-O '60. (MIRA 13:11)

1. Moskovskiy gosudarstvennyy universitet, kafedra neorganicheskoy
khimii.
(Beryllium compounds) (Naphthoic acid)

SEMENENKO, K.N.; LAVUT, Ye.A.

X-ray radiographic examination of sodium tellurite pentahydrate,
 $\text{Na}_2\text{TeO}_3 \cdot 5\text{H}_2\text{O}$. Vest. Mosk. un. Ser. 2: Khim. 15 no.6:27-29 N-D
'68. (MIRA 14:2)

1. Kafedra neorganicheskoy khimii Moskovskogo universiteta.
(Sodium tellurite--Spectra)

5.2100
AUTHORS:Kuvyrkin, O. N., Breysov, O. N.,
Novoselova, A. V., Semenenko, K. N.

688 50

S/076/60/034/02/012/044
B010/B015

TITLE:

On the Polymorphism of Beryllium Chloride 21

PERIODICAL:

Zhurnal fizicheskoy khimii, 1960, Vol 34, Nr 2, pp 343-348 (USSR)

ABSTRACT:

Beryllium chloride forms several polymorphous modifications. Since hitherto only the crystal structure of fibrous modifications has been investigated, the present study deals with the thermal and X-ray phase analysis of the polymorphism of beryllium chloride. The composition of the preparation applied is given (Table 1). Thermal structure analysis of this preparation was carried out with a PK-52 Kurnakov pyrometer and Pt/PtRh thermocouples. The Cu radiation of a BSV tube was used for the X-ray analyses, and the photographs were taken by RKD or RKU-86 cameras, and Unioam cameras at high temperatures, respectively. The results of the X-ray phase analyses are given (Tables 2-4). A rapid cooling-down of the beryllium chloride melt, or a crystallization from the gas phase, leads to a formation of the metastable α' -modification which is similar to silicon sulfide with respect to its structure. On heating the α' -modification is transformed at 250° into the cubic β' -modification which in turn is transformed into the stable

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On the Polymorphism of Beryllium Chloride

68850

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EO10/B015

β -modification at 340° . A slow cooling-down leads to the transformation melt $\xrightleftharpoons{425^{\circ}} \alpha(\text{BeCl}_2) \xrightleftharpoons{405^{\circ}} \beta\text{BeCl}_2$.
It is possible that the α -modification, the structure of which could not be identified, is the same as the α' -modification. The diagram obtained does, however, not comprise all polymorphous transformations of beryllium chloride, since N. V. Bychkov, e.g., (Ref 12) on crystallization of beryllium chloride in quartz vessels discovered a modification differing from the afore-mentioned modifications. There are 2 figures, 4 tables, and 12 references, 5 of which are Soviet.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet im. M. V. Lomonosova
(Moscow State University imeni M. V. Lomonosov)

SUBMITTED: April 24, 1958

Card 2/2

SEMENENKO, K.N.; KURDYUMOV, G.M.; GORDEYEV, I.V.

Heats of sublimation of beryllium oxysalts. Zhur.neorg.khim. 6
no.9:2025-2028 S '61. (MIRA 14:9)
(Beryllium salts)

SITDYKOVA, N.S.; TUROVA, N.Ya.; SEMENENKO, K.N.; NOVOSELOVA, A.V.

Compounds of beryllium chlorides with dialkyl sulfides. Zhur.
neorg.khim. 6 no.11:2512-2516 '61. (MIRA 14:10)
(Beryllium chloride) (Sulfide)

SEMENENKO, K.N.; KURGYUMOV, G.M.

Beryllium hydroxybenzoate. Zhur.neorg.khim. 6 no.11:2568-2571
'61. (MIRA 14:10)

(Beryllium compounds)

KURDYUMOV, G.M.; SEMENENKO, K.N.

Beryllium oxybenzoate compounds with aromatic hydrocarbons.
Zhur.neorg.khim. 7 no.9:2115-2121 S 1962. (MIRA 15:9)
(Beryllium compounds) (Hydrocarbons)

SEMENENKO, K.N.; KURDYUMOV, G.M.

Beryllium oxysalts of the alicyclic and aliphatic series. Zhur.neorg.khim.
7 no. 7:1512-1515 JI '62. (MIRA 16:3)

(Beryllium salts)

SEMIENENKO, K.N.; NAUMOVA, T.N.

Polymorphism of beryllium bromide and iodide. Zhur.strukt.khim.
4 no.1:67-72 Ja-F '63. (MIRA 16:2)

1. Moskovskiy gosudarstvennyy universitet imeni Lomonosova.
(Berillium bromide) (Berillium iodide)
(Polymorphism)

TUROVA, N.Ya.; SITDYKOVA, N.S.; NOVOSELOVA, A.V.; SEMENENKO, K.N.

Thermal decomposition of beryllium halide etherates. Zhur.neorg.-
khim. 8 no.2:528-531 F '63. (MIRA 16:5)

1. Moskovskiy gosudarstvennyy universitet imeni Lomonosova.
(Beryllium halides) (Ethers)

TUROVA, N.Ya.; SEMENENKO, K.N.; NOVOSELOVA, A.V.

Infrared absorption spectra and some problems of the complex formation of inorganic halides with ethers. Zhur.neorg.khim. 8 no.4:882-892
Ap '63. (MIRA 16:3)

1. Moskovskiy gosudarstvennyy universitet imeni Lomonosova.
(Halides) (Ethers) (Complex compounds--Absorption spectra)

MAKSIMOV, V.N.; SEMENENKO, K.N.

Thermal stability of cerium and neodymium acetates. Vest.Mosk.un.
Ser.2;Khim. 18 no.1:13-17 Ja-F '63. (MIRA 16:5)

1. Kafedra neorganicheskoy khimii Moskovskogo universiteta.
(Cerium acetates--Thermal properties)
(Neodymium acetates--Thermal properties)

SEMENENKO, K.N.; NAUMOVA, T.N.

Structure and some properties of aluminum halides. Zhur. neorg.
khim. 9 no.6:1316-1322 Je '63 (MIRA 17:8)

SEMENENKO, K.N.; TUROVA, N.Ya.

Structure of "onium" compounds of beryllium halides. Zhur.neorg.khim.
8 no.9:2093-2098 S '63. (MIRA 16:10)

TUROVA, N.Ya.; SEMENENKO, K.N.; NOVOSELOVA, A.V.

Compounds of beryllium chloride with phosphorus oxychloride.
Zhur.neorg.khim. 8 no.9:2155-2162 S '63. (MIRA 16:10)

1. Moskovskiy gosudarstvennyy universitet imeni Lomonosova.

KURDYUMOV, G.M.; SEMENENKO, K.N.

Heats of formation and the structure of the compounds of
beryllium hydroxybenzoate with benzene and styrene. Zhur.
neorg. khim. 8 no.11:2545-2548 N '63. (MIRA 17:1)

ACCESSION NR: AP4019502

S/0078/64/009/003/0765/0766

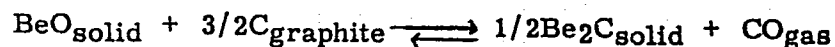
AUTHOR: Muratov, F. Sh. (Deceased); Semenenko, K. N.

TITLE: The cubic modification of beryllium oxide

SOURCE: Zhurnal neorg. khimii, v. 9, no. 3, 1964, 765-766

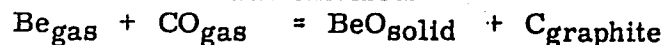
TOPIC TAGS: beryllium oxide, cubic modification, hexagonal modification, x ray data, phase transition

ABSTRACT: The phase transition of BeO from the hexagonal to the cubic modification at about 2000C and the existence of the cubic form only at temperatures above 2000C were reported by D. K. Smith, C. F. Cline, V. D. Frechette (J. Nucl. Mat., 6, 265 (1962)). An earlier work by F. Sh. Mauratov and A. V. Novoselova (Dokl. AN SSSR, 129, 334 (1959); 143, 599 (1962)), showed that in the reaction



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ACCESSION NR: AP4019502

the Be_2C dissociated and the Be was oxidized:

X-rays of the products showed hexagonal BeO, graphite, and a number of unidentified lines. By comparing these unidentified lines with data by Smith et al., it is established that they belong to the cubic modification of BeO. It is noted that in Muratov's work, the cubic form was observed in the sublimate at 1730C, was present simultaneously with the hexagonal form, and was stabilized by hardening. Orig. art. has: 1 table.

ASSOCIATION: Moskovskiy gosudarstvennyy universitet Khimicheskiy fakul'tet Kafedra neorganicheskoy khimii (Moscow State University, Chemistry Faculty, Department of Inorganic Chemistry).

SUBMITTED: 14Mar63

DATE ACQ: 31Mar64

ENCL: 00

SUB CODE: PH

NO REF SOV: 003

OTHER: 002

Card 2/2

TUROVA, N.Ya.; KEDROVA, N.S.; SEMENENKO, K.N.; NOVOSELOVA, A.V.

Interaction of etherates of beryllium chlorides and aluminum chlorides. Zhur.neorg.khim. 9 no.4:905-916 Ap '64.

(MIRA 17:4)

1. Moskovskiy gosudarstvennyy universitet imeni Lomonosova.

SEMENENKO, K.N.; TUROVA, N.Ya.

Structure of beryllium chloride hydrate-etherate. *Zhur. neorg. khim.*
9 no.4:1019-1020 Ap '64. (MIRA 17:4)

SEMENENKO, K.N.; NAUMOVA, T.N.; GOROKHOV, L.N.; SEMENOVA, G.A.; NOVOSELOVA, A.V.

Interaction between the chlorides of Al and Fe. Dokl. AN SSSR
154 no.1:169-170 Ja'64. (MIRA 17:2)

1. Moskovskiy gosudarstvennyy universitet im. M.V. Lomonosova.
2. Chlen-korrespondent AN SSSR (for Novoselova).

SEMENENKO, K.N.; NAUMOVA, T.N.; GOROKHOV, L.N.; NOVOSELOVA, A.V.

Interaction between the chlorides of aluminum and beryllium.
Dokl. AN SSSR 154 no. 3:648-649 Ja '64. (MIRA 17:5)

1. Moskovskiy gosudarstvennyy universitet im.M.V.Lomonosova.
2. Chlen-korrespondent AN SSSR (for Novoselcva).

SEMESENKO, K.N., TURKOVA, N.Ya.

Structure of beryllium chloride hydrate. Zhur. neorg. khim.
10 no.1:77-82 Ja '65. (MIR 18:11)

1. Moskovskiy gosudarstvennyy universitet imeni Lomonosova.
Submitted July 19, 1963.

SEMENENKO, K.N.; SUROV, V.N.

Study of sodium chloroberyllate and of the nature of its
reaction with beryllium chloride. Izv. AN SSSR. Neorg. mat.
1 no.11:1982-1989 N '65. (MIRA 18:12)

1. Khimicheskiy fakul'tet Moskovskogo gosudarstvennogo
universiteta imeni M.V. Lomonosova. Submitted June 11, 1965.

SEMENENKO, K.N.; GRIGOR'YEV, A.I.

Structure of potassium and sodium chloroberyllates.
Zhur.neorg.khim. 10 no.12:2591-2595 I '65.

(MIRA 19:1)

SEMENENKO, K.N.; TUROVA, N.Ya.

X-ray diffraction examination of a complex compound of
aluminum chloride with diethyl ether. Zhur.neorg.khim.
10 no.12:2830-2831 D '65.

(MIRA 19:1)

SEMEHENEO, K.N.; KEDROVA, N.S.; IOFA, B.Z.

Radiochemical study of sodium chloroberyllate and chloro-
aluminate. Zhur.neorg.khim. 10 no.12:2833-2834 D '65.
(MIRA 19±1)

NAMESTNIKOVA, L.N.; EDEL', Yu.P.; SEMENENKO, L.A.

Desirability of a postmortem quantitative determination of alcohol
in the contents and tissues of the stomach. Sud. - med. ekspert.
6 no.3255 J1-S'63. (MIRA 16810)

1. Kafedra sudebnoy meditsiny (zav. - N.P.Marchenko) Khar'-
kovskogo meditsinskogo instituta i Khar'kovskoye oblastnoye byuro
sudebno-meditsinskoy ekspertizy (nachal'nik V.M.Moiseyev).
(ALCOHOL IN THE BODY) (AUTOPSY)

MARCHENKO, N.P.; SEMENENKO, L.A.

Concerning the so-called "new sign" of intravital trauma proposed by V.I. Akopov. Sud.-med. ekspert. 2 no.3:56-58 J1-S '59.

(MIRA 13:4)

1. Kafedra sudebnoy meditsiny (zav. - prof. N.N. Bokarius) Khar'kovskogo meditsinskogo instituta i Khar'kovskoye oblastnoye byuro sudebnomeditsinskoy ekspertizy (nachal'nik N.P. Marchenko).
(WOUNDS)

SOTNIKOVA, L.L., dots.; SEMENENKO, L.A., sudebnomeditsinskiy ekspert (Khar'kov).

How did you dare? Zdorov'e 6 no.4:24 Ap '60.
(ABORTION)

(MIRA 13:8)

SEMNENKO, L.A.; POKUS, A.I.

Case of Poisoning by delagyl. Sud-med. ekspert. 6 nc.4:45
O-D'63 (MIRA 16:12)

1. Khar'kovskoye oblastnoye byuro sudebnomeditsinskoy eks-
pertizy (nachal'nik V.M.Moiseyev).

SEMENENKO, I. M.

Medicine

Personal hygiene; Kyiv, Med. vyd-vo URSR, 1951

Monthly List of Russian Accessions, Library of Congress, May 1952. UNCLASSIFIED

SEMENENKO, M.

Quality of vegetables and fruit. Sov. Horg. 35 no.5:21-22
My '62. (MIRA 15:5)

1. Starshiy tovaroved gorplodoovoshchtorga, g. Volgograd.
(Produce trade)

KAMYNIN, Yuliy Nikoalayeovich; MATVEYEV, M.G., kand. tekhn. nauk,
retsenzent; SEMENENKO, M.D., red.; STARODUB, T.A.,
tekhn. red. ~~tekhn. red.~~

[Contactless control diagrams in mine automation] Beekon-
taktnye skhemy upravleniia v shakhtnoi avtomatike. Kiev,
Gostekhizdat USSR, 1963. 214 p. (MIRA 17:1)

STARTSEV, V.I., otv. red.; ALEKSANDROV, B.S., red.; BELYAYEV, L.M., red.; BRUDZ', V.G., red.; VOYTOVETSKIY, V.K., red.; GALANIN, M.D., red.; DISTANOV, B.G., red.; KLIMOV, A.P., red.; SEMENENKO, M.G., red.; SHAMOVSKIY, L.M., red.

[Scintillators and scintillation materials] Stsintilliatory i stsintilliatsionnye materialy. Moskva, Gos. komitet Soveta Ministrov SSSR po khimii, 1960. 319 p. (MIRA 15:4)

1. Koordinatsionnoye soveshchaniye po stsintilliatoram. 2nd, 1957. (Scintillation counters)

CHERNAYEV, I.I.; ZHOLINOVSKAYA, N.N.; SEMENENKO, M.M.

Certain properties of tetraamino compounds of platinum(IV).
Zhur. neorg. khim. 6 no.1:44-47 '61. (MIRA 14:2)
(Platinum compounds)